

PATENT SPECIFICATION

(11) 1 549 845

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- (21) Application No. 14011/75 (22) Filed 4 April 1975
 (23) Complete Specification filed 25 March 1976
 (44) Complete Specification published 8 Aug. 1979
 (51) INT CL² C23C 9/02 9/04
 (52) Index at acceptance
 C7F 1B2 2A 2F 2M 2T 2Z2 2Z4 2Z8 4E 4F 4K 4X 5A
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(54) IMPROVEMENTS IN OR RELATING TO THE DIFFUSION COATING OF METAL OR OTHER ARTICLES

(71) I, THE SECRETARY OF STATE
 FOR DEFENCE, London, do hereby declare
 the invention, for which I pray that a patent
 may be granted to me, and the method by
 which it is to be performed, to be particu-
 larly described in and by the following state-
 ment:—

This invention relates to processes for coat-
 ing metal or other articles with diffusion coat-
 ings and particularly, though not exclusively,
 relates to coatings for gas turbine engine com-
 ponents, e.g. turbine blades, for increasing
 their high temperature corrosion resistance.

It is known to produce metallide or metal-
 lised coatings by constant pressure, halide
 activated, pack-cementation. Aluminised and
 chromised coatings have been applied to
 nickel-based, cobalt-based and iron-based sub-
 strates by this method.

In a known pack-cementation process such
 as is described in United States Patent
 3,257,230 to Wachtel, a method of coating
 cobalt-based or nickel-based alloys is disclosed
 in which a pack comprising a metallic coating
 material such as aluminium, chromium, iron
 or silicon, a carrier comprising a halide source
 such as ammonium halide, and a moderator
 metal are packed around the article to be
 coated and heated to between 760°C and
 1204°C for 15 minutes to 40 hours. During
 this heating, the aluminium or other coating
 material is transported to the surface of the
 article and deposited thereon.

In such a process, the transport of halide
 vapour between the coating material and the
 article is primarily by diffusion, and as a con-
 sequence such a process is fairly slow and has
 relatively poor 'throwing power', i.e. the trans-
 port by diffusion of the halide vapours will
 only occur effectively over very short distances.
 When coating articles on their external sur-
 faces, this poor 'throwing power' is usually
 not important, since by embedding the article
 to be coated in a particulate pack of the
 coating material the element which is to be
 transferred is in very close proximity of the
 article.

By contrast, the coating of internal surfaces
 such as bores is considerably more difficult to
 achieve by the pack cementation method. This
 is especially so in relation to the metallising
 of the surfaces of fine bores and cavities as are
 found for example in gas turbine engine
 components. United States Patent 3,079,276
 and British Patent 1,315,228 describe the fill-
 ing of holes and cavities of gas turbine engine
 components with a powder pack mix, but it is
 known that such practices are both difficult
 and time consuming to carry out and not
 always effective even for holes which are rela-
 tively large and of simple configuration.

It is also known to apply coatings of metals
 and compounds to internal surfaces by expos-
 ing the heated surfaces to a thermally decom-
 posable metal plating gas such as nickel car-
 bonyl. Such a process is described in British
 Patent 1,070,396.

The present invention provides an improved
 process for producing diffusion coatings on
 an article by metallizing or metallising and is
 particularly applicable to coating fine bores or
 narrow cavities in gas turbine engine com-
 ponents and other articles.

According to the present invention a pro-
 cess for coating a metal or other article with
 a diffusion coating comprises enclosing the
 article in a chamber together with a particu-
 late pack including coating material in ele-
 mental or chemically combined form, said
 coating material selected from the group com-
 prising aluminium, chromium, titanium, zir-
 conium, tantalum, niobium, yttrium, rare earth
 metals, boron and silicon, together with a
 halide activator, and cyclically varying the
 pressure of an inert gas or a reducing gas
 or a mixture of said gases within the chamber
 whilst maintaining the contents of the cham-
 ber at a temperature sufficient to transfer
 coating metal on to the surface of the article
 to form a diffusion coating thereon.

Preferably the method according to the
 invention is carried out at a pressure sub-
 stantially below atmospheric pressure that is,

below about 100 torr, and at a cycle frequency as high as is compatible with the transport of a sufficient quantity of the gas through the particulate pack per cycle. The ratio of upper pressure limit to lower pressure limit is preferably as high as is practicable and consistent with cycle frequency. Convenient pressure ranges are about 50 torr to about 10 torr, preferably with cycle frequencies of at least 2 cycles per minute. In general, higher frequencies are beneficial in increasing the ratio of coating thickness applied internally to that applied externally.

The halide activator is preferably selected from a group of inorganic halides having a low volatility at the coating temperature, such that halide loss from the chamber is low. Preferably, sufficient halide activator is used to ensure that some halide activator is retained in the pack at the end of the process. Examples of low volatility halide activators are shown in Table 1.

TABLE 1

Equilibrium Vapour, Sublimation or Dissociation Pressures of Some Low Volatility Halide Activators

Substance	Temp °C	Pressure torr	Substance	Temp °C	Pressure torr
AlF ₃	927	1.3	LiI	827	47
NaF	927	0.1	CrF ₃	785	0.01
NaCl	927	2.4	CrF ₂	927	approx 0.001
NaBr	927	4.8	CrCl ₂	927	approx 5.0
NaI	927	15.0	CrBr ₂	810	0.9
KF	927	1.2	CrI ₂	793	1.4
KCl	927	4.0	CoF ₂	927	0.05
KBr	927	6.5	FeF ₂	927	0.02
KI	927	12.3			

Activators of higher volatility are shown in Table 2.

TABLE 2

Equilibrium Vapour, Sublimation or Dissociation Pressures of Some High Volatility Activators

Substance	Temp °C	Pressure torr	Substance	Temp °C	Pressure torr
Cl ₂	-34	760	NH ₄ Br	397	760
Br ₂	61	760	AlCl ₃	180	760
I ₂	183	760	AlBr ₃	225	760
HCl	-167	760	AlI ₃	385	760
HBr	-35	760	FeCl ₂	319	760
HI	100	760	FeCl ₃	934	760
NH ₄ F	—	760	BeBr ₂	927	570
NH ₄ Cl	397	760			

The choice of halide activator must also take into account factors other than volatility. In particular, the activator must also be capable of entering into the necessary chemical equilibria which will lead to the to-and-fro gas transport and proper interaction with the coating material and the article to be coated, at the coating temperature which must not exceed the temperature at which significant degradation of the properties of the article occurs. In the case of certain nickel-base and cobalt-base alloys, for example, significant degradation can occur at coating temperatures above about 900°C. Other halides which may be used in the process of the invention are double halides for example sodium cryolite, Na₃AlF₆. It may also be desirable to employ a mixture of halide activators, for example NaF/NaCl/NaBr or KF/NaF/LiF to increase the efficiency of the deposition process.

When producing aluminised coatings by the method of the invention, aluminium fluoride, AlF₃, has been found to be particularly effective, good quality coatings of satisfactory uniformity and distribution having been obtained on both internal and external surfaces of nickel-base gas turbine blades. Preferably the proportions of the aluminium source material (coating material) and aluminium trifluoride are such that aluminium trifluoride crystals are not formed.

In one manner of operating the process in accordance with the invention, the article to be coated is kept out of physical contact with the particulate bed by placing the article to be coated inside a cage which is itself embedded within the particulate pack. A preferred construction for a cage is one that will permit vapours to pass from the particulate pack to the inside of the cage but which prevents or retards flow to the outside of the cage. One cage according to the invention has sides and an upper face of impermeable material, e.g. nickel sheet or plate, and the base of a mesh or gauze through which vapour can pass. This design has the effect of increasing the path length of the inert and/or reducing gas through the particulate bed, thereby promoting more effective entrainment of the active halide vapour.

The particulate pack may include a particulate filler such as a refractory oxide for support of the coating material or for dilution of the pack. A particulate filler comprising a refractory oxide may support a coating material such as liquid aluminium.

The articles coated by the process may be composed of any material that can be coated by pack cementation. Materials commonly coated by pack cementation are nickel-base, cobalt-base and iron-base alloys, and the refractory metals of Groups IV, V, and VI of the Periodic Table. In addition to these materials, carbon and carbon-containing materials, e.g. tungsten carbide, may be advantageously coated by the process.

An example of the process applied to aluminising and in accordance with the invention will now be described with reference to the drawing accompanying the Provisional Specification.

The drawing shows a leak-tight chamber and auxiliary plant in which the method may be carried out.

Referring to the drawing, the chamber includes a furnace tube 9 composed of mullite surrounded at its lower end by an alumina tube 5 surrounded in turn by an electrical heating element 4 and standing in a thermally insulated box 6 which has a nickel foil heat shield 7 on its upper surface. A gas turbine blade 1 of nickel-base alloy which is to be coated is located in a pack 2 comprising a powder mixture of aluminium, AlF_3 and Al_2O_3 powder retained in the furnace tube 9 by an aluminium disc 3.

The furnace tube 9 is connected by a pipe 20 to auxiliary equipment for continuously varying the pressure in the tube 9. The auxiliary equipment comprises a supply of argon 26 and a vacuum pump 27 connected to the pipe 20 by time controlled valves 24, 25 and needle valves 22, 23 respectively. A mercury manometer (not shown) is connected to a branch 21 of the pipe 20 and is used for

measuring the pressure fluctuations in the furnace tube 9.

The upper part of the furnace tube 9 is closed by an end plate 19 which is bolted to a flange 10 on the furnace tube. A pair of O-ring seals 13, 14 provide gas tight sealing between the end plate 19 and the furnace tube. A screw cap 28 engages with a thread on a cylindrical part of the end plate 19 and has an O-ring seal 12 which provides a gas tight seal between the end plate and the tube 17 which extends through the screw cap 28.

The upper part of the furnace tube 9 is water cooled, the water flowing through a copper pipe 18 to the upper end of the furnace tube. A stainless steel tube 17 surrounding the tube 18 carries the return water flow. The temperature of the pack is sensed by a thermocouple (not shown).

The method according to the invention is illustrated by the following examples:—

EXAMPLE 1

A gas turbine blade section in 'IN 100' alloy, bearing a hole of diameter about 1.5 mm and of length about 110 mm, was aluminised according to the method of the invention in a chamber. The method included embedding the blade section in a powder mix of 14 grams AlF_3 , 14 grams Al and 388 grams Al_2O_3 , raising the temperature of the chamber and its contents to 900°C and setting time-controlled valves to give a flow of argon into the chamber for 3 seconds as the pressure was increased from 6 torr to 28 torr, a constant pressure of about 28 torr for 20 seconds and then an exhaust period of 7 seconds to reduce the pressure to about 6 torr. After 10 hours at the same temperature, the chamber was cooled and the blade section removed. On examination, the surface of the hole was found to be uniformly coated with an aluminised layer of mean thickness about 35 μm . The thickness distribution of the coating along the length of the hole can be seen from the following figures:—

Distance from one end of hole (mm)	10	20	30	40	50
Coating thickness (μm)	40	40	35	30	30

EXAMPLE 2

In a further example, a turbine blade in 'IN 100' alloy bearing holes of diameter about 1.5 mm and of length about 70 mm, was aluminised for 5 hours at 900°C inside a nickel gauze cage which was itself embedded in a powder pack mix of 6.5 grams AlF_3 , 10.6 grams Al and 330 grams Al_2O_3 . The pressure range of argon was from 14 to 58 torr and the pressure cycle frequency was 6 cycles minute^{-1} . Bright metallic-looking and particularly smooth textured aluminised layers

were produced on both the internal and external surfaces of the blade. The layer thickness within the hole measured close to the top, mid-span and bottom was respectively 12, 8 and 12 μm . The layer thickness measured over the external surface at comparable positions was 25, 25 and 30 μm respectively.

EXAMPLE 3

In a further example a turbine blade in 'Nimonic 105' alloy (Nimonic being a Registered Trade Mark), bearing holes of two differing cross sections, about 0.8 and 1.5 mm, but of the same length about 60 mm, was aluminised for 6 hours at 900°C inside a nickel gauze cage which was itself embedded in a powder pack mix of 6.6 grams AlF_3 , 10.5 grams Al and 330 grams Al_2O_3 . The pressure range of argon was from 4 to 44 torr and the pressure cycle frequency was 6 cycles minute⁻¹. Aluminised layers of quality similar to those obtained in the preceding example were produced on both the internal and external surfaces of the blade. The internal surface of the larger cross section hole had thicknesses of 30, 20 and 30 μm respectively at the top, mid-span and bottom positions. The corresponding thicknesses for the smaller cross section hole was 15, 12 and 15 μm . External thicknesses at about the same positions were 65, 60 and 60 μm respectively.

EXAMPLE 4

In yet a further example of aluminising with AlF_3 , a turbine blade in 'MAR-M 246' alloy (MAR-M being a Registered Trade Mark), bearing holes of two different cross sections, one with major and minor axes of about 2 and 0.5 mm and the other of diameter about 2 mm, but both of length about 60 mm, were aluminised for 6 hours at 900°C inside a nickel gauze cage which was itself embedded in a powder pack mix of 6.6 grams AlF_3 , 10.6 grams Al and 330 grams Al_2O_3 . The pressure range of argon was from 12 to 52 torr and the pressure cycle frequency was 6 cycles minute⁻¹. Similar quality aluminised layers were produced on both the internal and external surfaces of the blade. The internal layer thickness along the length of the approximately elliptical section was 20, 15 and 25 μm close to the top, mid-span and bottom positions. The corresponding figures for the circular cross section were 40, 40 and 45 μm . Corresponding figures for the external surface were 60, 65 and 65 μm .

EXAMPLE 5

In an example of aluminising with NaCl as halide activator, a turbine blade in 'IN 100' alloy was aluminised for 5 hours at 900°C within a nickel gauze cage which was itself embedded in a powder pack mix of 20 grams NaCl, 14 grams Al and 300 grams Al_2O_3 . The pressure range of argon was from 8 torr

to 42 torr and the pressure cycle frequency was 6 cycles minute⁻¹. A layer of thickness 2 μm was produced within a hole of diameter 1.8 mm and of length 40 mm. The thickness of the external layer was about 16 μm .

EXAMPLE 6

In an example of aluminising with NaF as halide activator, an alloy section in 'IN 100' was aluminised for 5 hours at 900°C within a nickel gauze cage which was itself embedded in a powder pack mix of 14.7 grams NaF, 13.6 grams Al and 330 grams Al_2O_3 . The pressure range of argon was from 12 to 56 torr and the pressure cycle frequency was 6 cycles minute⁻¹. An aluminised layer of thickness 8 μm was produced within a hole of diameter about 1 mm and of length about 40 mm. The aluminised layer thickness over the external surface was about 30 μm .

The process of the invention may be used to apply various types of diffusion coatings to the internal and external surfaces of articles which hitherto have been applied by conventional pack cementation processes, e.g. aluminising, chromising, titanising, tantaliding, boronising and siliconising. To one skilled in the art, it will be appreciated that some activators may respond better than others to a particular metallising process, and some care must therefore be exercised in the choice of activators. Activators of low volatility that are not readily available may be synthesised within or outside the coating chamber prior to operation of the coating step and introduced into the pack. In one embodiment of the invention, SiCl_4 is synthesized from silicon and ammonium chloride, for use in siliconising.

WHAT I CLAIM IS:—

1. A process for coating a metal or other article with a diffusion coating comprising enclosing the article in a chamber, together with a particulate pack including coating material in elemental or chemically combined form said coating material selected from the group comprising aluminium, chromium, titanium, zirconium, tantalum, niobium, yttrium, rare earth metals, boron and silicon, together with a halide activator, and cyclically varying the pressure of an inert gas or a reducing gas or a mixture of said gases contained within the chamber whilst maintaining the contents of the chamber at a temperature sufficient to transfer coating material on to the surface of the article to form a diffusion coating thereon.

2. A process as claimed in Claim 1 in which the maximum pressure of the gas or gases is 100 torr.

3. A process as claimed in Claim 1 in which the maximum pressure of the gas or gases is 50 torr and the minimum pressure is 10 torr.

4. A process as claimed in any previous

- claim in which the pressure of the gas or gases is cyclically varied at a rate of at least 2 cycles per minute.
- 5 5. A process as claimed in any previous claim in which the halide activator has a low volatility at the temperature at which said diffusion coating is formed.
- 10 6. A process according to Claim 5 in which the coating material comprises aluminium and the halide activator comprises aluminium fluoride.
7. A process according to any previous claim in which the article is held out of contact with the particulate pack.
- 15 8. A process according to any previous claim in which the pack includes a particulate filler for supporting the coating material.
9. A process according to Claim 8 in which the carrier comprises a refractory oxide.
- 20 10. A process according to Claim 9 in which the refractory oxide comprises aluminium oxide.
11. A process according to any previous claim in which the halide activator is synthesised in the chamber.
- 25 12. Convert milligrams of sample/ml. of herein with reference to the Examples.
13. A metal article having a coating which has been coated onto said article by a process according to any previous claim.
- 30 14. An article composed of nickel-base alloy having a coating which has been coated onto said article by a process according to claim 6.
- 35 15. An article composed of iron-base alloy having a coating which has been coated onto said article by a process according to Claim 6.
- 40 16. An article composed of cobalt-base alloy having a coating which has been coated onto said article by a process according to Claim 6.
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Agent for the Applicant.

Printed for Her Majesty's Stationery Office, by the Courier Press, Leamington Spa, 1979
Published by The Patent Office, 25 Southampton Buildings, London, WC2A 1AY, from
which copies may be obtained.

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PROVISIONAL SPECIFICATION

1 SHEET

This drawing is a reproduction of
the Original on a reduced scale.

